

**LISTING OF THE CLAIMS**

This listing of the claims will replace all prior versions, and listings, of the claims in the application.

1. (Previously Presented) A method for treating a mammalian subject having a solid tumor *ex vivo*, comprising direct injection of a nucleic acid molecule encoding:
  - a) a polypeptide comprising the amino acid sequence of SEQ ID NO:2; or
  - b) a polypeptide comprising the amino acid sequence of SEQ ID NO:4;into cells of the tumor, such that the growth of the tumor is inhibited.
2. (Previously Presented) A method for modifying cells of a solid tumor *ex vivo* to express a B7-2 molecule comprising direct injection of a nucleic acid molecule encoding:
  - a) a polypeptide comprising the amino acid sequence of SEQ ID NO:2; or
  - b) a polypeptide comprising the amino acid sequence of SEQ ID NO:4;into cells of the tumor.
3. (Previously Presented) A method of increasing the immunogenicity of cells of a solid tumor *ex vivo* comprising direct injection of a nucleic acid molecule encoding:
  - a) a polypeptide comprising the amino acid sequence of SEQ ID NO:2; or
  - b) a polypeptide comprising the amino acid sequence of SEQ ID NO:4;into cells of the tumor, thereby increasing the immunogenicity of the tumor cells.
4. (Previously Presented) The method of any of claims 1-3, wherein the nucleic acid molecule encoding the polypeptide comprising the amino acid sequence of SEQ ID NO:2 or SEQ ID NO:4 comprises the nucleic acid sequence shown in SEQ ID NO:1 or SEQ ID NO:3.
5. (Canceled)
6. (Previously Presented) The method of any of claims 1-3, wherein the nucleic acid molecule encoding the polypeptide comprising the amino acid sequence of SEQ ID NO:2 or SEQ ID NO:4 is in a viral vector.

7. (Original) The method of claim 6, wherein the viral vector is selected from the group consisting of a retroviral vector, an adenoviral vector, and an adeno-associated viral vector.
8. (Previously Presented) The method of any of claims 1-3, wherein the nucleic acid molecule encoding the polypeptide comprising the amino acid sequence of SEQ ID NO:2 or SEQ ID NO:4 is a plasmid expression vector.
9. (Original) The method of any of claims 1-3, wherein the tumor cells are further transfected with at least one nucleic acid molecule encoding a B7-3 protein.
10. (Original) The method of any of claims 1-3, wherein the tumor cells are further injected with at least one nucleic acid molecule encoding at least one MHC class II  $\alpha$  chain protein and at least one MHC class II  $\beta$  chain protein in a form suitable for expression of the MHC class II  $\alpha$  chain protein(s) and the MHC class II  $\beta$  chain protein(s).
11. (Original) The method of any of claims 1-3, wherein the tumor cells are further injected with at least one nucleic acid molecule encoding at least one MHC class I  $\alpha$  chain protein in a form suitable for expression of the MHC class I protein(s).
12. (Original) The method of any of claims 1-3, wherein the tumor cells are further injected with a nucleic acid molecule encoding a  $\beta$ -2 microglobulin protein in a form suitable for expression of the  $\beta$ -2 microglobulin protein.
13. (Previously Presented) The method of any of claims 1-3, further comprising inhibiting expression of an MHC class II invariant chain in the tumor cells by transfection of the tumor cells with a nucleic acid molecule which is antisense to a regulatory or a coding region of the invariant chain gene.
14. (Original) The method of any of claims 1-3 wherein the solid tumor is selected from a group consisting of a carcinoma, sarcoma, melanoma and neuroblastoma.